Maine Newborn Screening Program

List of Conditions

Each baby born in Maine is screened for the conditions listed below. This list is correct as of ***July 1, 2008*** but may change as conditions are added to or removed from the testing panel. If you have any questions, please contact the Maine Newborn Screening Program at (207) 287-5357.

3-Hydroxy-3-methylglutaryl-CoA lyase deficiency
3-Methylcrotonyl-CoA carboxylase deficiency
Argininemia
Argininosuccinic acidemia
Beta-ketothiolase deficiency
Biotinidase deficiency
Carnitine palmitoyl transferase deficiency Type II
Carnitine uptake deficiency
Citrullinemia
Congenital adrenal hyperplasia
Congenital hypothyroidism
Cystic Fibrosis (CF)
Galactosemia
Glutaric acidemia type I
Glutaric acidemia type II
Homocystinuria
Hyperammonemia Hyperornithinemia Homocitrullinemia (HHH Syndrome)
Isovaleric acidemia
Long-chain acyl-CoA dehydrogenase (LCAD) deficiency
Long-chain hydroxyacyl-CoA dehydrogenase (LCHAD) deficiency
Maple syrup urine disease
Medium-chain acyl-CoA dehydrogenase (MCAD) deficiency
Methylmalonic acidemia
Multiple carboxylase deficiency
Phenylketonuria (PKU)
Propionic acidemia
Short-chain acyl-CoA dehydrogenase (SCAD) deficiency
Sickle cell disease/hemoglobin disorders
Trifunctional protein deficiency
Tyrosinemia type I
Tyrosinemia type II
Very long-chain acyl-CoA dehydrogenase (VLCAD) deficiency